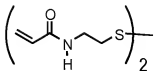


LISTING OF THE CLAIMS

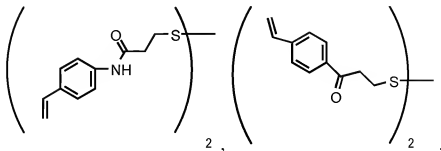
1. **(Previously Presented)** An antimicrobial lens comprising silver and a polymer comprising a monomer of Formula

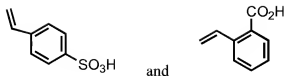


2. **(Canceled)**
3. **(Canceled)**
4. **(Previously Presented)** The antimicrobial lens of claim 1 wherein the lens is a soft contact lens.
5. **(Previously Presented)** The antimicrobial lens of claim 1 wherein the monomer of Formula I is present at about 0.01 to about 1.5 weight percent.
6. **(Previously Presented)** The antimicrobial lens of claim 1 wherein the monomer of Formula I is present at about 0.01 to about 0.8 weight percent.
7. **(Previously Presented)** The antimicrobial lens of claim 1 wherein the monomer of Formula I is present at about 0.01 to about 0.3 weight percent.
8. **(Previously Presented)** The antimicrobial lens of claim 1 wherein the monomer of Formula I is present at about 0.01 to about 0.2 weight percent.

9. **(Previously Presented)** The antimicrobial lens of claim 1 wherein the monomer of Formula I is present at about 0.01 to about 0.09 weight percent.
10. **(Previously Presented)** The antimicrobial lens of claim 1 wherein the lens is a silicone hydrogel.
11. **(Previously Presented)** The antimicrobial lens of claim 1 wherein, the lens is etafilcon A, balafilcon, A, aquafilcon A, lenefilcon A, or lotrafilcon A.
12. **(Canceled)**
13. **(Canceled)**
14. **(Previously Presented)** The antimicrobial lens of claim 1 wherein silver is present at about 20 ppm to about 1,200 ppm.
15. **(Previously Presented)** The antimicrobial lens of claim 1 wherein silver is present at about 20 ppm to about 600 ppm.
16. **(Previously Presented)** The antimicrobial lens of claim 1 wherein silver is present at about 20 ppm to about 150 ppm.
17. **(Previously Presented)** The antimicrobial lens of claim 1 wherein silver is present at about 20 ppm to about 75 ppm.
18. **(Canceled)**
19. **(Previously Presented)** The antimicrobial lens of claim 1 wherein silver is present at about 20 ppm to about 150 ppm and the monomer of Formula I is present at about 0.01 to about 1.5 weight percent.
20. **(Canceled)**

21. **(Previously Presented)** The antimicrobial lens of claim 11 wherein silver is present at about 20 ppm to about 150 ppm and the monomer of Formula I is present at about 0.01 to about 1.5 weight percent.
22. **(Original)** The antimicrobial lens of claim 21 wherein the lens is etafilcon A.
23. **(Original)** The antimicrobial lens of claim 21 wherein the lens is aquafilcon A.
24. **(Original)** The lens of claim 23 wherein silver is present at about 20 ppm to about 75 ppm.
25. **(Withdrawn)** The antimicrobial lens of claim 1 comprising a polymer comprising a monomer of Formula II.
26. **(Withdrawn)** The antimicrobial lens of claim 25 wherein,
a is 1-2,
 R^{11} is hydrogen or C_{1-3} alkyl,
 R^{12} is sulfonic acid, carboxylic acid, phosphonic acid, C_{1-6} alkyldisulfide,
 C_{1-6} alkylsulfide, phenyldisulfide, substituted phenyldisulfide or $NH-R^{13}$,
 R^{13} is thio C_{1-6} alkylcarbonyl.
27. **(Withdrawn)** The antimicrobial lens of claim 25 wherein the monomer of Formula II is selected from the group consisting of





28. **(Withdrawn)** The antimicrobial lens of claim 25 wherein the lens is a soft contact lens.
29. **(Withdrawn)** The antimicrobial lens of claim 25 wherein the monomer of Formula II is present at about 0.01 to about 1.5 weight percent.
30. **(Withdrawn)** The antimicrobial lens of claim 25 wherein the monomer of Formula II is present at about 0.01 to about 0.8 weight percent.
31. **(Withdrawn)** The antimicrobial lens of claim 25 wherein the monomer of Formula II is present at about 0.01 to about 0.3 weight percent.
32. **(Withdrawn)** The antimicrobial lens of claim 25 wherein the lens is etafilcon A, balafilcon, A, aquafilcon A, lenefilcon A, or lotrafilcon A.
33. **(Withdrawn)** The antimicrobial lens of claim 25 wherein silver is present at about 20 ppm to about 150 ppm and the monomer of Formula II is present at about 0.01 to about 1.5 weight percent.
34. **(Withdrawn)** The antimicrobial lens of claim 33 wherein the lens is etafilcon A or aquafilcon A.
35. **(Withdrawn)** The antimicrobial lens of claim 1 comprising a polymer comprising a monomer of Formula III.
36. **(Withdrawn)** The antimicrobial lens of claim 35 wherein,

p is 1-3;

b is 1-2;

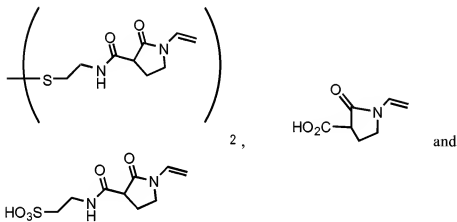
R²¹ is hydrogen;

R²² is sulfonic acid, phosphonic acid, carboxylic acid, thioC₁₋₆alkylcarbonyl, thioC₁₋₆alkylaminocarbonyl, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyldisulfide, substituted phenyldisulfide,

H₃OS-(CH₂)₁₋₆NHC(O) or

(HO)₂(O)P-(CH₂)₁₋₆NHC(O)-.

37. **(Withdrawn)** The antimicrobial lens of claim 35 wherein the monomer of Formula III is selected from the group consisting of

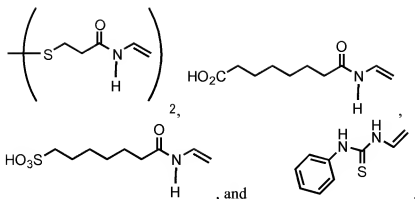


38. **(Withdrawn)** The antimicrobial lens of claim 35 wherein the lens is a soft contact lens.

39. **(Withdrawn)** The antimicrobial lens of claim 35 wherein the monomer of Formula III is present at about 0.01 to about 1.5 weight percent.

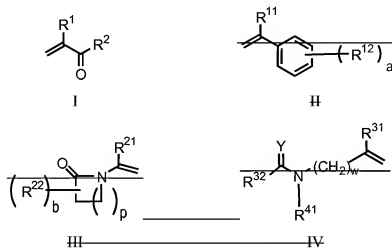
40. **(Withdrawn)** The antimicrobial lens of claim 35 wherein the monomer of Formula III is present at about 0.01 to about 0.8 weight percent.

41. **(Withdrawn)** The antimicrobial lens of claim 35 wherein the monomer of Formula III is present at about 0.01 to about 0.3 weight percent.
42. **(Withdrawn)** The antimicrobial lens of claim 35 wherein, the lens is etafilcon A, balafilcon, A, acquafilcon A, lenefilcon A, or lotrafilcon A.
43. **(Withdrawn)** The antimicrobial lens of claim 35 wherein silver is present at about 20 ppm to about 150 ppm and the monomer of Formula III is present at about 0.01 to about 1.5 weight percent.
44. **(Withdrawn)** The antimicrobial lens of claim 43 wherein the lens is etafilcon A or acquafilcon A.
45. **(Withdrawn)** The antimicrobial lens of claim 1 comprising a polymer comprising a monomer of Formula IV.
46. **(Withdrawn)** The antimicrobial lens of claim 45 wherein,
w is 0-1;
 R^{31} is hydrogen;
 R^{32} is amine, C_{1-3} alkylamine, phenylamine, substituted phenylamine;
thio C_{1-3} alkylcarbonyl;
 R^{41} is hydrogen.
47. **(Withdrawn)** The antimicrobial lens of claim 45 wherein the monomer of Formula IV is selected from the group consisting of



48. **(Withdrawn)** The antimicrobial lens of claim 45 wherein the lens is a soft contact lens.
49. **(Withdrawn)** The antimicrobial lens of claim 45 wherein the monomer of Formula IV is present at about 0.01 to about 1.5 weight percent.
50. **(Withdrawn)** The antimicrobial lens of claim 45 wherein the monomer of Formula IV is present at about 0.01 to about 0.8 weight percent.
51. **(Withdrawn)** The antimicrobial lens of claim 45 wherein the monomer of Formula IV is present at about 0.01 to about 0.3 weight percent.
52. **(Withdrawn)** The antimicrobial lens of claim 45 wherein the lens is etafilcon A, balafilcon, A, acquafilcon A, lenefilcon A, or lotrafilcon A.
53. **(Withdrawn)** The antimicrobial lens of claim 45 wherein silver is present at about 20 ppm to about 150 ppm and the monomer of Formula IV is present at about 0.01 to about 1.5 weight percent.
54. **(Withdrawn)** The antimicrobial lens of claim 53 wherein the lens is etafilcon A or acquafilcon A.

55. **(Currently Amended)** A method of producing an antimicrobial lens comprising, up to about 200 ppm silver and a polymer comprising a monomer of Formula I, II, III or IV



wherein

R¹ is hydrogen or C₁₋₆alkyl;

R^2 is $-OR^3$, $-NH-R^3$, $-S-(CH_2)_d-R^3$, or $-(CH_2)_d-R^3$, wherein

d is 0-8;

R³ is substituted C₁₋₆alkyl

where the alkyl substituents are selected from one or more members of the group consisting of carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, nitrile, thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyldisulfide, urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea, phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted phenyldisulfide, substituted C₁₋₆alkylurea, substituted phenylurea, substituted C₁₋₆alkylthiourea, and substituted phenylthiourea

wherein the C₁₋₆alkyldisulfide, phenyldisulfide, C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and phenylthiourea substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl,

carboxylic acid, sulfonic acid, phosphonic acid, amine,
amidine, acetamide, and nitrile;

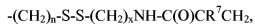


wherein R^4 , R^5 , and R^6 are independently selected from the
group consisting of hydrogen, halogen, hydroxyl, and

C_{1-6} alkyl,

q is 1-6, and

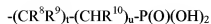
m is 0-6;



wherein R^7 is hydrogen or C_{1-6} alkyl,

n is 1-6, and

x is 1-6;



wherein R^8 , R^9 , and R^{10} are independently selected from the
group consisting of hydrogen, halogen, hydroxyl, and

C_{1-6} alkyl,

t is 1-6, and

u is 0-6;

phenyl;

benzyl;

pyridinyl;

pyrimidinyl;

pyrazinyl;

benzimidazolyl;

benzothiazolyl;

benzotriazolyl;

naphthaloyl;

quinolinyl;

indolyl;

thiadiazolyl;

triazolyl;

4-methylpiperidin-1-yl;
4-methylpiperazin-1-yl;
substituted phenyl;
substituted benzyl;
substituted pyridinyl;
substituted pyrimidinyl;
substituted pyrazinyl;
substituted benzimidazolyl;
substituted benzothiazolyl;
substituted benzotriazolyl;
substituted naphthaloyl;
substituted quinolinyl;
substituted indolyl;
substituted thiadiazolyl;
substituted triazolyl;
substituted 4-methylpiperidin-1-yl; or
substituted 4-methylpiperazin-1-yl,

wherein the substituents are selected from one or more members of the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, sulfonic acid, phosphonic acid, hydroxyl, carboxylic acid, amine, amidine, N-(2-aminopyrimidine)sulfonyl, N-(aminopyridine)sulfonyl, N-(aminopyrazine)sulfonyl, N-(2-aminopyrimidine)carbonyl, N-(aminopyridine)carbonyl, N-(aminopyrazine)carbonyl, N-(2-aminopyrimidine)phosphonyl, N-(2-aminopyridine)phosphonyl, N-(aminopyrazine)phosphonyl, N-(aminobenzimidazolyl)sulfonyl, N-(aminobenzothiazolyl)sulfonyl, N-(aminobenzotriazolyl)sulfonyl, N-(aminoindolyl)sulfonyl, N-(aminothiazolyl)sulfonyl,

N-(aminotriazolyl)sulfonyl,
 N-(amino-4-methylpiperidinyl)sulfonyl,
 N-(amino-4-methylpiperazinyl)sulfonyl,
 N-(aminobenzimidazolyl)carbonyl,
 N-(aminobenzothiazolyl)carbonyl,
 N-(aminobenzotriazolyl)carbonyl, N-(aminoindolyl)carbonyl,
 N-(aminothiazolyl)carbonyl,
 N-(aminotriazolyl)carbonyl,
 N-(amino-4-methylpiperidinyl)carbonyl,
 N-(amino-4-methylpiperazinyl)carbonyl,
 N-(2-aminobenzimidazolyl)phosphonyl,
 N-(2-aminobenzothiazolyl)phosphonyl,
 N-(2-aminobenzotriazolyl)phosphonyl,
 N-(2-aminoindolyl)phosphonyl,
 N-(2-aminothiazolyl)phosphonyl,
 N-(2-aminotriazolyl)phosphonyl,
 N-(amino-4-methylpiperidinyl) phosphonyl,
 N-(amino-4-methylpiperazinyl) phosphonyl, acetamide, nitrile,
 thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyl disulfide,
 urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea,
 phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted
 phenyldisulfide, substituted C₁₋₆alkylurea, substituted
 C₁₋₆alkylthiourea, substituted phenylurea, and substituted
 phenylthiourea

wherein the C₁₋₆alkyldisulfide, phenyldisulfide,
 C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and
 phenylthiourea substituents are selected from the group
 consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl,
 carboxylic acid, sulfonic acid, phosphonic acid, amine,
 amidine, acetamide, and nitrile;

a is 1-5;

R^{+1} is hydrogen or C_{1-6} alkyl;

R^{+2} is hydroxyl, sulfonic acid, phosphonic acid, carboxylic acid, acetamide, thio C_{1-6} alkylearbonyl, C_{1-6} alkyldisulfide, C_{1-6} alkylsulfide, phenyl disulfide, urea, C_{1-6} alkylurea, phenylurea, thiourea, C_{1-6} alkylthiourea, phenylthiourea, $-OR^{+3}$, $-NH-R^{+3}$, $-S-(CH_2)_d-R^{+3}$, $-(CH_2)_d-R^{+3}$, $-C(O)NH-(CH_2)_d-R^{+3}$, $-C(O)-(CH_2)_d-R^{+3}$; substituted C_{1-6} alkyldisulfide, substituted phenyldisulfide, substituted C_{1-6} alkylurea, substituted phenylurea, substituted phenylthiourea or substituted C_{1-6} alkylthiourea wherein the substituents are selected from the group consisting of C_{1-6} alkyl, halo C_{1-6} alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;

where

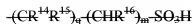
d is 0-8;

R^{+3} is thio C_{1-6} alkylearbonyl;

—— substituted C_{1-6} alkyl

where the alkyl substituents are selected from one or more members of the group consisting of hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, nitrile, thiol, C_{1-6} alkyldisulfide, C_{1-6} alkylsulfide, phenyldisulfide, urea, C_{1-6} alkylurea, phenylurea, thiourea, C_{1-6} alkylthiourea, phenylthiourea, substituted C_{1-6} alkyldisulfide, substituted phenyldisulfide, substituted C_{1-6} alkylurea, substituted phenylurea, substituted C_{1-6} alkylthiourea and substituted phenylthiourea

wherein the C_{1-6} alkyldisulfide, phenyldisulfide, C_{1-6} alkylurea, C_{1-6} alkylthiourea, phenylurea, and phenylthiourea substituents are selected from the group consisting of C_{1-6} alkyl, halo C_{1-6} alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;

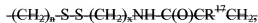


where R^{14} , R^{15} , and R^{16} are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and

C_{1-6} alkyl;

q is 1-6, and

m is 0-6;



where R^{17} is hydrogen or C_{1-6} alkyl;

n is 1-6, and

x is 1-6;



where R^{18} , R^{19} , and R^{20} are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and

C_{1-6} alkyl;

t is 1-6, and

u is 0-6;

phenyl;

benzyl;

pyridinyl;

pyrimidinyl;

pyrazinyl;

benzimidazolyl;

benzothiazolyl;

benzotriazolyl;

naphthaloyl;

quinolinyl;

indolyl;

thiadiazolyl;

triazolyl;

4-methylpiperidin-1-yl;

4-methylpiperazin-1-yl;

substituted phenyl;

substituted-benzyl;
 substituted-pyridinyl;
 substituted-pyrimidinyl;
 substituted-pyrazinyl;
 substituted-benzimidazolyl;
 substituted-benzothiazolyl;
 substituted-benzotriazolyl;
 substituted-naphthaloyl;
 substituted-quinolinyl;
 substituted-indolyl;
 substituted-thiadiazolyl;
 substituted-triazolyl;
 substituted-4-methylpiperidin-1-yl; or
 substituted-4-methylpiperazin-1-yl

wherein the substituents are selected from one or more
 members of the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl,
 halogen, sulfonic acid, phosphonic acid, hydroxyl, carboxylic
 acid, amine, amidine, N-(2-aminopyrimidine)sulfonyl,
 N-(aminopyridine)sulfonyl, N-(aminopyrazine)sulfonyl,
 N-(2-aminopyrimidine)carbonyl, N-(aminopyridine)carbonyl,
 N-(aminopyrazine)carbonyl,
 N-(2-aminopyrimidine)phosphonyl,
 N-(2-aminopyridine)phosphonyl,
 N-(aminopyrazine)phosphonyl,
 N-(aminobenzimidazolyl)sulfonyl,
 N-(aminobenzothiazolyl)sulfonyl,
 N-(aminobenzotriazolyl)sulfonyl, N-(aminoindolyl)sulfonyl,
 N-(aminothiazolyl)sulfonyl,
 N-(aminotriazolyl)sulfonyl,
 N-(amino-4-methylpiperidinyl)sulfonyl,
 N-(amino-4-methylpiperazinyl)sulfonyl,

N-(aminobenzimidazolyl)carbonyl;
 N-(aminobenzothiazolyl)carbonyl;
 N-(aminobenzotriazolyl)carbonyl, N-(aminoindolyl)carbonyl;
 N-(aminothiazolyl)carbonyl;
 N-(aminotriazolyl)carbonyl;
 N-(amino-4-methylpiperidinyl)carbonyl;
 N-(amino-4-methylpiperazinyl)carbonyl;
 N-(2-aminobenzimidazolyl)phosphonyl;
 N-(2-aminobenzothiazolyl)phosphonyl;
 N-(2-aminobenzotriazolyl)phosphonyl;
 N-(2-aminoindolyl)phosphonyl;
 N-(2-aminothiazolyl)phosphonyl;
 N-(2-aminotriazolyl)phosphonyl;
 N-(amino-4-methylpiperidinyl)phosphonyl;
 N-(amino-4-methylpiperazinyl)phosphonyl; acetamide, nitrile,
 thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyl disulfide,
 urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea,
 phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted
 phenyldisulfide, substituted C₁₋₆alkylurea, substituted
 C₁₋₆alkylthiourea, substituted phenylurea, and substituted
 phenylthiourea

wherein the C₁₋₆alkyldisulfide, phenyldisulfide,
 C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and
 phenylthiourea substituents are selected from the group
 consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl,
 carboxylic acid, sulfonic acid, phosphonic acid, amine,
 amidine, acetamide, and nitrile;

b is 1-5;

p is 1-5;

R²¹ is hydrogen;

R^{22} is hydroxyl, sulfonic acid, phosphonic acid, carboxylic acid, thio C_{1-6} alkylearbonyl, thio C_{1-6} alkylaminocarbonyl, C_{1-6} alkyldisulfide, phenyldisulfide, $-C(O)NH(CH_2)_{1-6}-SO_3H$, $-C(O)NH(CH_2)_{1-6}-P(O)(OH)_{2-3}-OR^{23}$, $-NH-R^{23}$, $-C(O)NH(CH_2)_d-R^{23}$, $-S(CH_2)_d-R^{23}$, $-(CH_2)_d-R^{23}$, urea, C_{1-6} alkylurea, phenylurea, thiourea, C_{1-6} alkylthiourea, phenylthiourea, substituted C_{1-6} alkyldisulfide, substituted phenyldisulfide, substituted C_{1-6} alkylurea, substituted, C_{1-6} alkylthiourea-substituted phenylurea or substituted phenylthiourea wherein the substituents are selected from the group consisting of C_{1-6} alkyl, halo C_{1-6} alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile,

where

d is 0-8;

R^{23} is—thio C_{1-6} alkylearbonyl,

C_{1-6} alkyl,

substituted C_{1-6} alkyl

where the alkyl substituents are selected from one or more members of the group consisting of C_{1-6} alkyl, halo C_{1-6} alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, nitrile, thiol, C_{1-6} alkyldisulfide, C_{1-6} alkylsulfide, phenyldisulfide, urea, C_{1-6} alkylurea, phenylurea, thiourea, C_{1-6} alkylthiourea, phenylthiourea, substituted C_{1-6} alkyldisulfide, substituted phenyldisulfide, substituted C_{1-6} alkylurea, substituted phenylurea, substituted C_{1-6} alkylthiourea, and substituted phenylthiourea

wherein the C_{1-6} alkyldisulfide, phenyldisulfide, C_{1-6} alkylurea, C_{1-6} alkylthiourea, phenylurea, and phenylthiourea substituents are selected from the group consisting of C_{1-6} alkyl, halo C_{1-6} alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;



where R^{24} , R^{25} , and R^{26} are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and

C_{1-6} alkyl;

q is 1-6, and

m is 0-6



where R^{27} is hydrogen or C_{1-6} alkyl;

n is 1-6, and

x is 1-6;



where R^{28} , R^{29} , and R^{30} are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and

C_{1-6} alkyl;

t is 1-6, and

u is 0-6;

phenyl;

benzyl;

pyridinyl;

pyrimidinyl;

pyrazinyl;

benzimidazolyl;

benzothiazolyl;

benzotriazolyl;

naphthaloyl;

quinolinyl;

indolyl;

thiadiazolyl;

triazolyl;

4-methylpiperidin-1-yl;

4-methylpiperazin-1-yl;

substituted-phenyl;
substituted-benzyl;
substituted-pyridinyl;
substituted-pyrimidinyl;
substituted-pyrazinyl;
substituted-benzimidazolyl;
substituted-benzothiazolyl;
substituted-benzotriazolyl;
substituted-naphthaloyl;
substituted-quinolinyl;
substituted-indolyl;
substituted-thiadiazolyl;
substituted-triazolyl;
substituted 4-methylpiperidin-1-yl; or
substituted 4-methylpiperazin-1-yl;

wherein the substituents are selected from one or more members of the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, sulfonic acid, phosphonic acid, hydroxyl, carboxylic acid, amine, amidine, N-(2-aminopyrimidine)sulfonyl, N-(aminopyridine)sulfonyl, N-(aminopyrazine)sulfonyl, N-(2-aminopyrimidine)carbonyl, N-(aminopyridine)carbonyl, N-(aminopyrazine)carbonyl, N-(2-aminopyrimidine)phosphonyl, N-(2-aminopyridine)phosphonyl, N-(aminopyrazine)phosphonyl, N-(aminobenzimidazolyl)sulfonyl, N-(aminobenzothiazolyl)sulfonyl, N-(aminobenzotriazolyl)sulfonyl, N-(aminoindolyl)sulfonyl, N-(aminothiazolyl)sulfonyl, N-(aminotriazolyl)sulfonyl, N-(amino-4-methylpiperidinyl)sulfonyl,

N-(amino-4-methylpiperazinyl)sulfonyl,
 N-(aminobenzimidazolyl)carbonyl,
 N-(aminobenzothiazolyl)carbonyl,
 N-(aminobenzotriazolyl)carbonyl, N-(aminoindolyl)carbonyl,
 N-(aminothiazolyl)carbonyl,
 N-(aminotriazolyl)carbonyl,
 N-(amino-4-methylpiperidinyl)carbonyl,
 N-(amino-4-methylpiperazinyl)carbonyl,
 N-(2-aminobenzimidazolyl)phosphonyl,
 N-(2-aminobenzothiazolyl)phosphonyl,
 N-(2-aminobenzotriazolyl)phosphonyl,
 N-(2-aminoindolyl)phosphonyl,
 N-(2-aminothiazolyl)phosphonyl,
 N-(2-aminotriazolyl)phosphonyl,
 N-(amino-4-methylpiperidinyl)phosphonyl,
 N-(amino-4-methylpiperazinyl)phosphonyl, acetamide, nitrile,
 thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyl disulfide,
 urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea,
 phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted
 phenyldisulfide, substituted C₁₋₆alkylurea, substituted
 C₁₋₆alkylthiourea, substituted phenylurea, and substituted
 phenylthiourea

wherein the C₁₋₆alkyldisulfide, phenyldisulfide,
 C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and
 phenylthiourea substituents are selected from the group
 consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl,
 carboxylic acid, sulfonic acid, phosphonic acid, amine,
 amidine, acetamide, and nitrile;

w is 0-1;

Y is oxygen or sulfur;

R²¹ is hydrogen or C₁₋₆alkyl;

R^{32} is hydroxyl, sulfonic acid, phosphonic acid, carboxylic acid, thio C_{1-6} alkylecarbonyl, thio C_{1-6} alkylaminocarbonyl, $C(O)NH-(CH_2)_d-R^{33}$, $-O-R^{33}$, $-NH-R^{33}$, $-S-(CH_2)_d-R^{33}$, $-(CH_2)_d-R^{33}$, C_{1-6} alkyldisulfide, phenyldisulfide, urea, C_{1-6} alkylurea, phenylurea, thiourea, C_{1-6} alkylthiourea, phenylthiourea, C_{1-6} alkylamine, phenylamine, substituted C_{1-6} alkyldisulfide, substituted phenyldisulfide, substituted phenylurea, substituted C_{1-6} alkylamine, substituted phenylamine, substituted phenylthiourea, substituted C_{1-6} alkylurea or substituted C_{1-6} alkylthiourea wherein the substituents are selected from the group consisting of C_{1-6} alkyl, halo C_{1-6} alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile

where

d is 0-8;

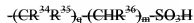
R^{32} is thio C_{1-6} alkylecarbonyl,

C_{1-6} alkyl,

substituted C_{1-6} alkyl

where the alkyl substituents are selected from one or more members of the group consisting of C_{1-6} alkyl, halo C_{1-6} alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, nitrile, thiol, C_{1-6} alkyldisulfide, C_{1-6} alkylsulfide, phenyldisulfide, urea, C_{1-6} alkylurea, phenylurea, thiourea, C_{1-6} alkylthiourea, phenylthiourea, substituted C_{1-6} alkyldisulfide, substituted phenyldisulfide, substituted C_{1-6} alkylurea, substituted phenylurea, substituted C_{1-6} alkylthiourea or substituted phenylthiourea wherein the C_{1-6} alkyldisulfide, phenyldisulfide, C_{1-6} alkylurea, C_{1-6} alkylthiourea, phenylurea, and phenylthiourea substituents are selected from the group consisting of C_{1-6} alkyl, halo C_{1-6} alkyl, halogen, hydroxyl,

carboxylic acid, sulfonic acid, phosphonic acid, amine,
amidine, acetamide, and nitrile;



where R^{34} , R^{35} , and R^{36} are independently selected from the
group consisting of hydrogen, halogen, hydroxyl, and

C_{1-6} alkyl;

q is 1-6, and

m is 0-6;



where R^{37} is hydrogen or C_{1-6} alkyl;

n is 1-6, and

x is 1-6;



where R^{38} , R^{39} , and R^{40} are independently selected from the
group consisting of hydrogen, halogen, hydroxyl, and

C_{1-6} alkyl;

t is 1-6, and

u is 0-6;

phenyl;

benzyl;

pyridinyl;

pyrimidinyl;

pyrazinyl;

benzimidazolyl;

benzothiazolyl;

benzotriazolyl;

naphthaloyl;

quinolinyl;

indolyl;

thiadiazolyl;

triazolyl;

4-methylpiperidin-1-yl;
4-methylpiperazin-1-yl;
substituted-phenyl;
substituted-benzyl;
substituted-pyridinyl;
substituted-pyrimidinyl;
substituted-pyrazinyl;
substituted-benzimidazolyl;
substituted-benzothiazolyl;
substituted-benzotriazolyl;
substituted-naphthaloyl;
substituted-quinolinyl;
substituted-indolyl;
substituted-thiadiazolyl;
substituted-triazolyl;
substituted 4-methylpiperidin-1-yl; or
substituted 4-methylpiperazin-1-yl;

wherein the substituents are selected from one or more members of the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, sulfonic acid, phosphonic acid, hydroxyl, carboxylic acid, amine, amidine, N-(2-aminopyrimidine)sulfonyl, N-(aminopyridine)sulfonyl, N-(aminopyrazine)sulfonyl, N-(2-aminopyrimidine)carbonyl, N-(aminopyridine)carbonyl, N-(aminopyrazine)carbonyl, N-(2-aminopyrimidine)phosphonyl, N-(2-aminopyridine)phosphonyl, N-(aminopyrazine)phosphonyl, N-(aminobenzimidazolyl)sulfonyl, N-(aminobenzothiazolyl)sulfonyl, N-(aminobenzotriazolyl)sulfonyl, N-(aminoindolyl)sulfonyl, N-(aminothiazolyl)sulfonyl,

N-(aminotriazolyl)sulfonyl,
 N-(amino-4-methylpiperidinyl)sulfonyl,
 N-(amino-4-methylpiperazinyl)sulfonyl,
 N-(aminobenzimidazolyl)carbonyl,
 N-(aminobenzothiazolyl)carbonyl,
 N-(aminobenzotriazolyl)carbonyl, N-(aminoindolyl)carbonyl,
 N-(aminothiazolyl)carbonyl,
 N-(aminotriazolyl)carbonyl,
 N-(amino-4-methylpiperidinyl)carbonyl,
 N-(amino-4-methylpiperazinyl)carbonyl,
 N-(2-aminobenzimidazolyl)phosphonyl,
 N-(2-aminobenzothiazolyl)phosphonyl,
 N-(2-aminobenzotriazolyl)phosphonyl,
 N-(2-aminoindolyl)phosphonyl,
 N-(2-aminothiazolyl)phosphonyl,
 N-(2-aminotriazolyl)phosphonyl,
 N-(amino-4-methylpiperidinyl)phosphonyl,
 N-(amino-4-methylpiperazinyl)phosphonyl, acetamide, nitrile,
 thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyl disulfide,
 urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea,
 phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted
 phenyldisulfide, substituted C₁₋₆alkylurea, substituted
 C₁₋₆alkylthiourea, substituted phenylurea, and substituted
 phenylthiourea

wherein the C₁₋₆alkyldisulfide, phenyldisulfide,
 C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and
 phenylthiourea substituents are selected from the group
 consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl,
 carboxylic acid, sulfonic acid, phosphonic acid, amine,
 amidine, acetamide, and nitrile;

R⁴⁴ is hydrogen, C₁₋₆alkyl, phenyl, C₁₋₆alkylearbonyl, phenylearbonyl, substituted C₁₋₆alkyl, substituted phenyl, substituted C₁₋₆alkylearbonyl or substituted phenylearbonyl,

— wherein

— the substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile.

where the method comprises the steps of

- (a) preparing a lens comprising a monomer of Formula I, ~~II, III or IV~~ and
- (b) treating said lens with a silver solution.

56. **(Original)** The method of claim 55 wherein the silver solution is aqueous silver nitrate having a concentration of about 0.1 µg/mL to about .3 g/mL.

57. **(Currently Amended)** The method of claim 55 wherein, treating comprises soaking the lens comprising a polymer of a monomer of Formula I, ~~II, III or IV~~ with a silver solution.

58. **(Currently Amended)** The method of claim 55 wherein, the lens comprising a polymer of a monomer of Formula I, ~~II, III or IV~~ is soaking for about 2 minutes to about 2 hours.

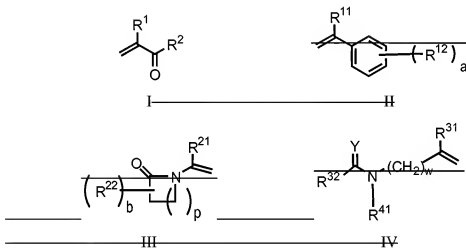
59. **(Currently Amended)** The method of claim 55 wherein, treating comprises storing the lens comprising a polymer of a monomer of Formula I, ~~II, III or IV~~ with a silver solution for about 20 minutes to about 5 years.

60. **(Currently Amended)** An antimicrobial lens comprising up to about 200 ppm silver and a polymer comprising a binding monomer wherein said antimicrobial lens can reversibly bind silver.

61. **(Original)** The antimicrobial lens of claim 60 wherein the binding monomer has a stability constant of about 2 to about 7.3.

62. **(Canceled)**

63. **(Currently Amended)** A method of reducing the adverse effects associated with microbial production in the eye of a mammal comprising providing an antimicrobial lens, wherein said lens comprises, up to about 200 ppm silver and a polymer comprising a monomer of the Formula I, ~~II, III or IV~~



wherein

R^1 is hydrogen or C_{1-6} alkyl;

R^2 is $-OR^3$, $-NH-R^3$, $-S-(CH_2)_d-R^3$, or $-(CH_2)_d-R^3$, wherein

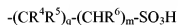
d is 0-8;

R^3 is substituted C_{1-6} alkyl

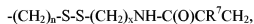
where the alkyl substituents are selected from one or more members of the group consisting of carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, nitrile, thiol, C_{1-6} alkyldisulfide, C_{1-6} alkylsulfide, phenyldisulfide, urea, C_{1-6} alkylurea, phenylurea, thiourea, C_{1-6} alkylthiourea, phenylthiourea, substituted C_{1-6} alkyldisulfide, substituted

phenyldisulfide, substituted C₁₋₆alkylurea, substituted phenylurea, substituted C₁₋₆alkylthiourea, and substituted phenylthiourea

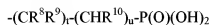
wherein the C₁₋₆alkyldisulfide, phenyldisulfide, C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and phenylthiourea substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;



wherein R⁴, R⁵, and R⁶ are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and C₁₋₆alkyl,
q is 1-6, and
m is 0-6;



wherein R⁷ is hydrogen or C₁₋₆alkyl,
n is 1-6, and
x is 1-6;



wherein R⁸, R⁹, and R¹⁰ are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and C₁₋₆alkyl,
t is 1-6, and
u is 0-6;

phenyl;
benzyl;
pyridinyl;
pyrimidinyl;
pyrazinyl;
benzimidazolyl;

benzothiazolyl;
benzotriazolyl;
naphthaloyl;
quinolinyl;
indolyl;
thiadiazolyl;
triazolyl;
4-methylpiperidin-1-yl;
4-methylpiperazin-1-yl;
substituted phenyl;
substituted benzyl;
substituted pyridinyl;
substituted pyrimidinyl;
substituted pyrazinyl;
substituted benzimidazolyl;
substituted benzothiazolyl;
substituted benzotriazolyl;
substituted naphthaloyl;
substituted quinolinyl;
substituted indolyl;
substituted thiadiazolyl;
substituted triazolyl;
substituted 4-methylpiperidin-1-yl; or
substituted 4-methylpiperazin-1-yl,

wherein the substituents are selected from one or more members of the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, sulfonic acid, phosphonic acid, hydroxyl, carboxylic acid, amine, amidine, N-(2-aminopyrimidine)sulfonyl, N-(aminopyridine)sulfonyl, N-(aminopyrazine)sulfonyl, N-(2-aminopyrimidine)carbonyl, N-(aminopyridine)carbonyl, N-(aminopyrazine)carbonyl,

N-(2-aminopyrimidine)phosphonyl,
 N-(2-aminopyridine)phosphonyl,
 N-(aminopyrazine)phosphonyl,
 N-(aminobenzimidazolyl)sulfonyl,
 N-(aminobenzothiazolyl)sulfonyl,
 N-(aminobenzotriazolyl)sulfonyl, N-(aminoindolyl)sulfonyl,
 N-(aminothiazolyl)sulfonyl,
 N-(aminotriazolyl)sulfonyl,
 N-(amino-4-methylpiperidinyl)sulfonyl,
 N-(amino-4-methylpiperazinyl)sulfonyl,
 N-(aminobenzimidazolyl)carbonyl,
 N-(aminobenzothiazolyl)carbonyl,
 N-(aminobenzotriazolyl)carbonyl, N-(aminoindolyl)carbonyl,
 N-(aminothiazolyl)carbonyl,
 N-(aminotriazolyl)carbonyl,
 N-(amino-4-methylpiperidinyl)carbonyl,
 N-(amino-4-methylpiperazinyl)carbonyl,
 N-(2-aminobenzimidazolyl)phosphonyl,
 N-(2-aminobenzothiazolyl)phosphonyl,
 N-(2-aminobenzotriazolyl)phosphonyl,
 N-(2-aminoindolyl)phosphonyl,
 N-(2-aminothiazolyl)phosphonyl,
 N-(2-aminotriazolyl)phosphonyl,
 N-(amino-4-methylpiperidinyl) phosphonyl,
 N-(amino-4-methylpiperazinyl) phosphonyl, acetamide, nitrile,
 thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyl disulfide,
 urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea,
 phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted
 phenyldisulfide, substituted C₁₋₆alkylurea, substituted
 C₁₋₆alkylthiourea, substituted phenylurea, and substituted
 phenylthiourea

wherein the C₁₋₆alkyldisulfide, phenyldisulfide, C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and phenylthiourea substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;

a is 1-5;

R¹¹ is hydrogen or C₁₋₆alkyl;

R¹² is hydroxyl, sulfonic acid, phosphonic acid, carboxylic acid, acetamide, thioC₁₋₆alkylearbonyl, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyl disulfide, urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea, phenylthiourea, -OR¹³, -NH-R¹³, -S-(CH₂)_d-R¹³, -(CH₂)_d-R¹³, -C(O)NH-(CH₂)_d-R¹³, -C(O)-(CH₂)_d-R¹³; substituted C₁₋₆alkyldisulfide, substituted phenyldisulfide, substituted C₁₋₆alkylurea, substituted phenylurea, substituted phenylthiourea or substituted C₁₋₆alkylthiourea wherein the substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;

where

d is 0-8;

R¹³ is thioC₁₋₆alkylearbonyl;

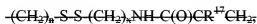
—— substituted C₁₋₆alkyl

where the alkyl substituents are selected from one or more members of the group consisting of hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, nitrile, thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyldisulfide, urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea, phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted phenyldisulfide, substituted C₁₋₆alkylurea, substituted phenylurea, substituted C₁₋₆alkylthiourea and substituted phenylthiourea

wherein the C₁₋₆alkyldisulfide, phenyldisulfide, C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and phenylthiourea substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;



where R¹⁴, R¹⁵, and R¹⁶ are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and C₁₋₆alkyl;
q is 1-6, and
m is 0-6;



where R¹⁷ is hydrogen or C₁₋₆alkyl;
n is 1-6, and
x is 1-6;



where R¹⁸, R¹⁹, and R²⁰ are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and C₁₋₆alkyl;
t is 1-6, and
u is 0-6;

phenyl;
benzyl;
pyridinyl;
pyrimidinyl;
pyrazinyl;
benzimidazolyl;
benzothiazolyl;
benzotriazolyl;
naphthaloyl;

quinolinyl;
 indolyl;
 thiadiazolyl;
 triazolyl;
 4-methylpiperidin-1-yl;
 4-methylpiperazin-1-yl;
 substituted-phenyl;
 substituted-benzyl;
 substituted-pyridinyl;
 substituted-pyrimidinyl;
 substituted-pyrazinyl;
 substituted-benzimidazolyl;
 substituted-benzothiazolyl;
 substituted-benzotriazolyl;
 substituted-naphthaloyl;
 substituted-quinolinyl;
 substituted-indolyl;
 substituted-thiadiazolyl;
 substituted-triazolyl;
 substituted-4-methylpiperidin-1-yl; or
 substituted-4-methylpiperazin-1-yl

wherein the substituents are selected from one or more
 members of the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl,
 halogen, sulfonic acid, phosphonic acid, hydroxyl, carboxylic
 acid, amine, amidine, N-(2-aminopyrimidine)sulfonyl,
 N-(aminopyridine)sulfonyl, N-(aminopyrazine)sulfonyl,
 N-(2-aminopyrimidine)carbonyl, N-(aminopyridine)carbonyl,
 N-(aminopyrazine)carbonyl,
 N-(2-aminopyrimidine)phosphonyl,
 N-(2-aminopyridine)phosphonyl,
 N-(aminopyrazine)phosphonyl,

N-(aminobenzimidazolyl)sulfonyl,
 N-(aminobenzothiazolyl)sulfonyl,
 N-(aminobenzotriazolyl)sulfonyl, N-(aminoindolyl)sulfonyl,
 N-(aminothiazolyl)sulfonyl,
 N-(aminotriazolyl)sulfonyl,
 N-(amino-4-methylpiperidinyl)sulfonyl,
 N-(amino-4-methylpiperazinyl)sulfonyl,
 N-(aminobenzimidazolyl)carbonyl,
 N-(aminobenzothiazolyl)carbonyl,
 N-(aminobenzotriazolyl)carbonyl, N-(aminoindolyl)carbonyl,
 N-(aminothiazolyl)carbonyl,
 N-(aminotriazolyl)carbonyl,
 N-(amino-4-methylpiperidinyl)carbonyl,
 N-(amino-4-methylpiperazinyl)carbonyl,
 N-(2-aminobenzimidazolyl)phosphonyl,
 N-(2-aminobenzothiazolyl)phosphonyl,
 N-(2-aminobenzotriazolyl)phosphonyl,
 N-(2-aminoindolyl)phosphonyl,
 N-(2-aminothiazolyl)phosphonyl,
 N-(2-aminotriazolyl)phosphonyl,
 N-(amino-4-methylpiperidinyl)phosphonyl,
 N-(amino-4-methylpiperazinyl)phosphonyl, acetamide, nitrile,
 thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyl disulfide,
 urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea,
 phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted
 phenyldisulfide, substituted C₁₋₆alkylurea, substituted
 C₁₋₆alkylthiourea, substituted phenylurea, and substituted
 phenylthiourea
 wherein the C₁₋₆alkyldisulfide, phenyldisulfide,
 C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and
 phenylthiourea substituents are selected from the group

consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;

b is 1-5;

p is 1-5;

R²¹ is hydrogen;

R²² is hydroxyl, sulfonic acid, phosphonic acid, carboxylic acid, thioC₁₋₆alkylcarbonyl, thioC₁₋₆alkylaminocarbonyl, C₁₋₆alkyldisulfide, phenyldisulfide, C(O)NH(CH₂)₄₋₆SO₃H, C(O)NH(CH₂)₄₋₆P(O)(OH)₂, OR²³, -NH-R²³, C(O)NH(CH₂)_d-R²², -(CH₂)_d-R²², urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea, phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted phenyldisulfide, substituted C₁₋₆alkylurea, substituted, C₁₋₆alkylthiourea-substituted phenylurea or substituted phenylthiourea wherein the substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile,

where

d is 0-8;

R²³ is thioC₁₋₆alkylcarbonyl,

C₁₋₆alkyl,

substituted C₁₋₆alkyl

where the alkyl substituents are selected from one or more members of the group consisting of C₁₋₆alkyl, halo C₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, nitrile, thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyldisulfide, urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea, phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted phenyldisulfide, substituted C₁₋₆alkylurea, substituted phenylurea, substituted C₁₋₆alkylthiourea, and substituted phenylthiourea

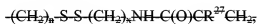
wherein the C₁₋₆alkyldisulfide, phenyldisulfide, C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and phenylthiourea substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;



where R²⁴, R²⁵, and R²⁶ are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and C₁₋₆alkyl;

q is 1-6, and

m is 0-6



where R²⁷ is hydrogen or C₁₋₆alkyl;

n is 1-6, and

x is 1-6;



where R²⁸, R²⁹, and R²⁰ are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and C₁₋₆alkyl;

t is 1-6, and

u is 0-6;

phenyl;

benzyl;

pyridinyl;

pyrimidinyl;

pyrazinyl;

benzimidazolyl;

benzothiazolyl;

benzotriazolyl;

naphthaloyl;

quinolinyl;
indolyl;
thiadiazolyl;
triazolyl;
4-methylpiperidin-1-yl;
4-methylpiperazin-1-yl;
substituted-phenyl;
substituted-benzyl;
substituted-pyridinyl;
substituted-pyrimidinyl;
substituted-pyrazinyl;
substituted-benzimidazolyl;
substituted-benzothiazolyl;
substituted-benzotriazolyl;
substituted-naphthaloyl;
substituted-quinolinyl;
substituted-indolyl;
substituted-thiadiazolyl;
substituted-triazolyl;
substituted-4-methylpiperidin-1-yl; or
substituted-4-methylpiperazin-1-yl;

wherein the substituents are selected from one or more members of the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, sulfonic acid, phosphonic acid, hydroxyl, carboxylic acid, amine, amidine, N-(2-aminopyrimidine)sulfonyl, N-(aminopyridine)sulfonyl, N-(aminopyrazine)sulfonyl, N-(2-aminopyrimidine)carbonyl, N-(aminopyridine)carbonyl, N-(aminopyrazine)carbonyl, N-(2-aminopyrimidine)phosphonyl, N-(2-aminopyridine)phosphonyl, N-(aminopyrazine)phosphonyl,

N-(aminobenzimidazolyl)sulfonyl,
 N-(aminobenzothiazolyl)sulfonyl,
 N-(aminobenzotriazolyl)sulfonyl, N-(aminoindolyl)sulfonyl,
 N-(aminothiazolyl)sulfonyl,
 N-(aminotriazolyl)sulfonyl,
 N-(amino-4-methylpiperidinyl)sulfonyl,
 N-(amino-4-methylpiperazinyl)sulfonyl,
 N-(aminobenzimidazolyl)carbonyl,
 N-(aminobenzothiazolyl)carbonyl,
 N-(aminobenzotriazolyl)carbonyl, N-(aminoindolyl)carbonyl,
 N-(aminothiazolyl)carbonyl,
 N-(aminotriazolyl)carbonyl,
 N-(amino-4-methylpiperidinyl)carbonyl,
 N-(amino-4-methylpiperazinyl)carbonyl,
 N-(2-aminobenzimidazolyl)phosphonyl,
 N-(2-aminobenzothiazolyl)phosphonyl,
 N-(2-aminobenzotriazolyl)phosphonyl,
 N-(2-aminoindolyl)phosphonyl,
 N-(2-aminothiazolyl)phosphonyl,
 N-(2-aminotriazolyl)phosphonyl,
 N-(amino-4-methylpiperidinyl)phosphonyl,
 N-(amino-4-methylpiperazinyl)phosphonyl, acetamide, nitrile,
 thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyl disulfide,
 urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea,
 phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted
 phenyldisulfide, substituted C₁₋₆alkylurea, substituted
 C₁₋₆alkylthiourea, substituted phenylurea, and substituted
 phenylthiourea
 wherein the C₁₋₆alkyldisulfide, phenyldisulfide,
 C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and
 phenylthiourea substituents are selected from the group

consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;

w is 0-1;

Y is oxygen or sulfur;

R²¹ is hydrogen or C₁₋₆alkyl;

R²² is hydroxyl, sulfonic acid, phosphonic acid, carboxylic acid, thioC₁₋₆alkylcarbonyl, thioC₁₋₆alkylaminocarbonyl, C(O)NH-(CH₂)_d-R²³, -O-R²³, -NH-R²³, -S-(CH₂)_d-R²³, -(CH₂)_d-R²³, C₁₋₆alkyldisulfide, phenyldisulfide, urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea, phenylthiourea, C₁₋₆alkylamine, phenylamine, substituted C₁₋₆alkyldisulfide, substituted phenyldisulfide, substituted phenylurea, substituted C₁₋₆alkylamine, substituted phenylamine, substituted phenylthiourea, substituted C₁₋₆alkylurea or substituted C₁₋₆alkylthiourea wherein the substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile

where

d is 0-8;

R²³ is thioC₁₋₆alkylcarbonyl,

C₁₋₆alkyl,

substituted C₁₋₆alkyl

where the alkyl substituents are selected from one or more members of the group consisting of C₁₋₆alkyl, halo C₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, nitrile, thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyldisulfide, urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea, phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted phenyldisulfide, substituted C₁₋₆alkylurea, substituted phenylurea,

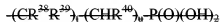
substituted C_{1-6} alkylthiourea or substituted phenylthiourea
 wherein the C_{1-6} alkyldisulfide, phenyldisulfide, C_{1-6} alkylurea, C_{1-6} alkylthiourea, phenylurea, and phenylthiourea substituents are selected from the group consisting of C_{1-6} alkyl, halo C_{1-6} alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;



where R^{24} , R^{25} , and R^{26} are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and C_{1-6} alkyl;
 q is 1-6, and
 m is 0-6;



where R^{27} is hydrogen or C_{1-6} alkyl;
 n is 1-6, and
 x is 1-6;



where R^{28} , R^{30} , and R^{40} are independently selected from the group consisting of hydrogen, halogen, hydroxyl, and C_{1-6} alkyl;
 t is 1-6, and
 u is 0-6;

phenyl;
 benzyl;
 pyridinyl;
 pyrimidinyl;
 pyrazinyl;
 benzimidazolyl;
 benzothiazolyl;

benzotriazolyl;
naphthaloyl;
quinolinyl;
indolyl;
thiadiazolyl;
triazolyl;
4-methylpiperidin-1-yl;
4-methylpiperazin-1-yl;
substituted-phenyl;
substituted-benzyl;
substituted-pyridinyl;
substituted-pyrimidinyl;
substituted-pyrazinyl;
substituted-benzimidazolyl;
substituted-benzothiazolyl;
substituted-benzotriazolyl;
substituted-naphthaloyl;
substituted-quinolinyl;
substituted-indolyl;
substituted-thiadiazolyl;
substituted-triazolyl;
substituted-4-methylpiperidin-1-yl; or
substituted-4-methylpiperazin-1-yl;

wherein the substituents are selected from one or more members of the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, sulfonic acid, phosphonic acid, hydroxyl, carboxylic acid, amine, amidine, N-(2-aminopyrimidine)sulfonyl, N-(aminopyridine)sulfonyl, N-(aminopyrazine)sulfonyl, N-(2-aminopyrimidine)carbonyl, N-(aminopyridine)carbonyl, N-(aminopyrazine)carbonyl, N-(2-aminopyrimidine)phosphonyl,

N-(2-aminopyridine)phosphonyl,
 N-(aminopyrazine)phosphonyl,
 N-(aminobenzimidazolyl)sulfonyl,
 N-(aminobenzothiazolyl)sulfonyl,
 N-(aminobenzotriazolyl)sulfonyl, N-(aminoindolyl)sulfonyl,
 N-(aminothiazolyl)sulfonyl,
 N-(aminotriazolyl)sulfonyl,
 N-(amino-4-methylpiperidinyl)sulfonyl,
 N-(amino-4-methylpiperazinyl)sulfonyl,
 N-(aminobenzimidazolyl)carbonyl,
 N-(aminobenzothiazolyl)carbonyl,
 N-(aminobenzotriazolyl)carbonyl, N-(aminoindolyl)carbonyl,
 N-(aminothiazolyl)carbonyl,
 N-(aminotriazolyl)carbonyl,
 N-(amino-4-methylpiperidinyl)carbonyl,
 N-(amino-4-methylpiperazinyl)carbonyl,
 N-(2-aminobenzimidazolyl)phosphonyl,
 N-(2-aminobenzothiazolyl)phosphonyl,
 N-(2-aminobenzotriazolyl)phosphonyl,
 N-(2-aminoindolyl)phosphonyl,
 N-(2-aminothiazolyl)phosphonyl,
 N-(2-aminotriazolyl)phosphonyl,
 N-(amino-4-methylpiperidinyl)phosphonyl,
 N-(amino-4-methylpiperazinyl)phosphonyl, acetamide, nitrile,
 thiol, C₁₋₆alkyldisulfide, C₁₋₆alkylsulfide, phenyl disulfide,
 urea, C₁₋₆alkylurea, phenylurea, thiourea, C₁₋₆alkylthiourea,
 phenylthiourea, substituted C₁₋₆alkyldisulfide, substituted
 phenyldisulfide, substituted C₁₋₆alkylurea, substituted
 C₁₋₆alkylthiourea, substituted phenylurea, and substituted
 phenylthiourea

wherein the C₁₋₆alkyldisulfide, phenyldisulfide, C₁₋₆alkylurea, C₁₋₆alkylthiourea, phenylurea, and phenylthiourea substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile;

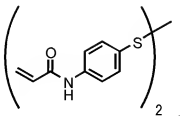
R⁴⁺ is hydrogen, C₁₋₆alkyl, phenyl, C₁₋₆alkylcarbonyl, phenylcarbonyl, substituted C₁₋₆alkyl, substituted phenyl, substituted C₁₋₆alkylcarbonyl or substituted phenylcarbonyl,

— wherein

— the substituents are selected from the group consisting of C₁₋₆alkyl, haloC₁₋₆alkyl, halogen, hydroxyl, carboxylic acid, sulfonic acid, phosphonic acid, amine, amidine, acetamide, and nitrile.

64. **(Currently Amended)** An antimicrobial lens comprising up to about 200 ppm silver, wherein said lens has sufficient movement on the eye of a patient.
65. **(Currently Amended)** The lens of claim 64 wherein said lens has sufficient movement on between having about 50 to about 100 percent movement of patients wearing said lens.
66. **(Currently Amended)** The lens of claim 64 wherein said lens has sufficient movement on between having about 75 to about 100 percent movement of patients wearing said lens.
67. **(Currently Amended)** The lens of claim 64 wherein said lens has sufficient movement on between having about 90 to about 100 percent movement of patients wearing said lens.
68. **(Currently Amended)** An antimicrobial lens comprising up to about 200 ppm silver, wherein said lens inhibits microbial production by at least 25%.

69. **(Original)** The lens of claim 68 wherein said lens inhibits microbial production by at least about 50% to at least about 99%.
70. **(Original)** The lens of claim 68 wherein said lens inhibits microbial production by at least about 80% to at least about 99%.
71. **(Currently Amended)** An antimicrobial lens comprising up to about 200 ppm silver, wherein said lens has sufficient movement on the eye of a patient and said lens inhibits microbial production by at least 25%.
72. **(Currently Amended)** The lens of claim 71 wherein said lens has sufficient movement on between ~~having~~ about 50% to about 100% of patients wearing said ~~lens movement~~ and said lens inhibits microbial production by 75% to about 100%.
73. **(Original)** The lens of claim 1 wherein said silver reversibly binds to said monomer.
74. **(Previously Presented)** An antimicrobial lens comprising silver and a polymer comprising a monomer of Formula



75. **(Previously Presented)** An antimicrobial lens comprising silver and a polymer comprising a monomer of Formula

